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In the Claims

Please replace all prior versions, and listings, of claims in the application with the following list of claims:

1-91. (Canceled)

- 92. (Currently Amended) A method for the production of transgenic animals comprising:
- a) transfecting a first non-human differentiated somatic cell or cell-line with a transgene construct containing a first-DNA sequence;
- b) selecting a transfected cell or cell-line into which <u>thesaid first</u> DNA sequence has been inserted into <u>theits</u> genome <u>of said first non-human differentiated somatic cell or cell-line</u>, <u>wherein the DNA sequence is operably linked to a promoter</u>;
- c) performing a first nuclear transfer procedure to generate a first transgenic animal at least heterozygous for thesaid first DNA sequence, wherein the first transgenic animal and the selected transfected cell or cell-line are of the same species;
- d) performing a biopsy or other cell selection technique to obtain cells to establish a second non-human differentiated somatic cell or cell-line from thesaid first transgenic animal;
- e) characterizing <u>thesaid</u> second non-human differentiated somatic cell or cell-line <u>using</u> molecular biology methods to ensure that the second non-human differentiated somatic cell or cell-line is at least heterozygous for <u>thesaid first-DNA</u> sequence; and
- f) performing a second nuclear transfer procedure with at least one one-cell of <u>thesaid</u> second non-human differentiated somatic cell or cell-line to produce at least a second transgenic animal at least heterozygous for said first DNA sequence; and,
- g) producing <u>athe</u> second transgenic animal <u>at least heterozygous for the DNA sequence</u>, <u>wherein the second transgenic animal and the second non-human differentiated somatic cell or cell-line are of the same species.</u>
- 93. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> first transgenic animal is at an embryonic stage of development.

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94. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> first transgenic animal is at a fetal stage of development.

- 95. (Currently Amended) The method of claim 92, further comprising developing <u>thesaid</u> first transgenic animal into an adult non-human animal.
- 96. (Currently Amended) The method of claim 92, wherein thesaid first transgenic animal is a mammal.
- 97. (Currently Amended) The method of claim 92, wherein thesaid first DNA sequence encodes a desired protein.
- 98. (Currently Amended) The method of claim 92, wherein the genetic composition of the said first transgenic animal is characterized to confirm the presence and expression of the DNA sequence transgene.
- 99. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> first nuclear transfer procedure further comprises transferring the nucleus of <u>thesaid</u> transfected cell into a suitable enucleated recipient cell of the same species, thereby obtaining a reconstituted cell.
- 100. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> first transgenic animal is biopsied so as to characterize the genome of <u>thesaid</u> first transgenic animal.
- 101. (Currently Amended) The method of claim 92, wherein at least one cell from thesaid second non-human differentiated somatic cell or cell-line is expanded through cell culture techniques prior tofor use in thesaid second round of nuclear transfer so as to allow for the production of produce a multiplicity of animals transgenic for thesaid DNA sequence of interest.
- 102. (Currently Amended) The method of claim 96, wherein the source of <u>thesaid</u> differentiated somatic cell or cell-line is an ungulate.

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103. (Currently Amended) The method of claim 102, wherein <u>thesaid</u> ungulate is selected from the group consisting of bovine, ovine, porcine, equine, caprine and buffalo.

- 104. (Currently Amended) A method of preparing a genetically engineered transgenic mammal, comprising:
- (a) inseminating a first female non-human mammal recipient with semen from a transgenic non-human animal of the same species known to have that has a transgene present and expressed;
 - (b) obtaining a transgenic non-human embryo from thesaid first female recipient;
 - (c) obtaining a <u>differentiated</u> somatic cell from <u>thesaid</u> embryo;
- (d) culturing thesaid differentiated somatic cell in a suitable medium, such that a differentiated somatic cell line is obtained; and,
- (e) performing a nuclear transfer procedure with <u>cells from thesaid</u> differentiated somatic <u>cell lineeells</u> to produce at least one transgenic mammal at least heterozygous for <u>thesaid</u> transgene, wherein <u>thesaid</u> transgene encodes a desired gene actuated by a tissue-specific promoter, wherein the transgenic mammal is of the same species as the differentiated somatic cell line; and,
 - (f) producing the transgenic mammal.
- 105. (Canceled)
- 106. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> second non-human differentiated somatic cell or cell-line is obtained from an embryonic goat on or after day 10 of embryogenesis.
- 107. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> second non-human differentiated somatic cell or cell line is kept in an airtight container.

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108. (Currently Amended) The method of claim 92, wherein thesaid first-DNA sequence codes for a biopharmaceutical protein product.

109. (Currently Amended) The method of claim [[108]]92, wherein the promoter is asaid first DNA sequence encodes a desired gene that is actuated by at least one beta-casein promoter.

110. (Canceled)

- 111. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> second non-human differentiated somatic cell or cell-line is obtained from <u>thesaid</u> first transgenic animal by tissue dissociation means including enzymatic means and/or mechanical means.
- 112. (Currently Amended) The method of claim 92, wherein thesaid second non-human differentiated somatic cell is a fibroblast, cumulus cell, neural cell, mammary cell or a myocyte or thesaid second non-human differentiated somatic cell-line is from a fibroblast, cumulus cell, neural cell, mammary cell or a myocyte.

113. (Canceled)

- 114. (Currently Amended) The method of claim 104, wherein <u>thesaid</u> transgene codes for a biopharmaceutical protein product.
- 115. (Currently Amended) The method of claim [[114]]104, wherein thesaid tissue-specific promoter is a beta-case in promoter.

116. (Canceled)

117. (Currently Amended) The method of claim 104, wherein <u>thesaid</u> second non-human differentiated somatic cell or cell-line is obtained from <u>thesaid</u> first transgenic animal by tissue dissociation means including enzymatic means and/or mechanical means.

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118. (Currently Amended) The method of claim 104, wherein <u>thesaid</u> second non-human differentiated somatic cell is a fibroblast, cumulus cell, neural cell, mammary cell or a myocyte or <u>thesaid</u> second non-human differentiated somatic cell-line is from a fibroblast, cumulus cell, neural cell, mammary cell or a myocyte.

- 119. (Currently Amended) The method of claim 92, wherein the DNAsaid transgene construct comprises a nucleic acid sequence encodesencoding a human polypeptide.
- 120. (Currently Amended) The method of claim 92, wherein <u>insertion into the genome of the DNA sequence results insaid transgene construct is capable of knocking out the expression of a genean endogenous gene of theto said first transgenic animal.</u>
- 121. (Currently Amended) The method of claim 119, wherein the <u>DNA sequencesaid</u> transgene construct further comprises a promoter, and wherein the <u>nucleic acidDNA sequence</u> is under the control of <u>thesaid</u> promoter.
- 122. (Currently Amended) The method of claim 121, wherein <u>thesaid</u> promoter is a tissue_specific promoter.
- 123. (Currently Amended) The method of claim 122, wherein <u>thesaid</u> tissue-specific promoter <u>is a promoter preferentially expressed induces expression</u> in mammary gland epithelial cells.
- 124. (Currently Amended) The method of claim 123, wherein <u>thesaid</u> promoter is selected from the group consisting of a beta-casein promoter, beta-lactoglobin promoter, whey acid protein promoter and lactalbumin promoter.
- 125. (Currently Amended) The method of claim 121, wherein <u>thesaid</u> promoter is a caprine promoter.

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126. (Currently Amended) The method of claim 119, wherein thesaid nucleic acid encodes a polypeptide is selected from the group consisting of a hormone, an immunoglobin, a plasma protein, and an enzyme.

- 127. (Currently Amended) The method of claim 119, wherein thesaid nucleic acid encodes a polypeptide is selected from the group consisting of an alpha-1 proteinase inhibitor, an alkaline phosphotase, an angiogenin, an extracellular superoxide dismutase, a fibrogen, a glucocerebrosidase, a glutamate decarboxylase, a human serum albumin, a myelin basis protein, a proinsulin, a soluble CD4, a lactoferrin, a lactoglobulin, a lysozyme, a lactoalbumin, an erythropoietin, a tissue plasminogen activator, a human growth factor, an antithrombin III, an insulin, a prolactin, and an alpha-1-antitrypsin.
- 128. (Currently Amended) The method of claim 92, wherein <u>thesaid</u> second non-human differentiated somatic cell is a fibroblast or <u>thesaid</u> second non-human differentiated somatic cell-line is from a fibroblast.
- 129. (Currently Amended) The method of claim 128, wherein <u>thesaid</u> fibroblast is a primary fibroblast.
- 130. (Currently Amended) The method of claim 128, wherein <u>thesaid</u> fibroblast is a primary derived fibroblast.